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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/462,962	06/25/2001	STEPHEN PHILIP JACKSON	MEWE-010	5713
24353 7:	590 01/11/2005		EXAMINER	
	FIELD & FRANCIS	ROBINSON, HOPE A		
1900 UNIVERSITY AVE SUITE 200			ART UNIT	PAPER NUMBER
EAST PALO ALTO, CA 94303			1653	

DATE MAILED: 01/11/2005

Please find below and/or attached an Office communication concerning this application or proceeding.

		Applicati n N .	Applicant(s)		
		09/462,962	JACKSON ET AL.		
	Office Action Summary	Examin r	Art Unit		
		Hope A. Robinson	1653		
Period fo	The MAILING DATE of this communication app or Reply	pears on the cover sheet with the c	orrespondence address		
THE - Exte after - If the - If NO - Failu Any	ORTENED STATUTORY PERIOD FOR REPLY MAILING DATE OF THIS COMMUNICATION. nsions of time may be available under the provisions of 37 CFR 1.13 SIX (6) MONTHS from the mailing date of this communication. e period for reply specified above is less than thirty (30) days, a reply period for reply is specified above, the maximum statutory period vere to reply within the set or extended period for reply will, by statute, reply received by the Office later than three months after the mailing ed patent term adjustment. See 37 CFR 1.704(b).	36(a). In no event, however, may a reply be time y within the statutory minimum of thirty (30) days will apply and will expire SIX (6) MONTHS from the application to become ABANDONE!	nely filed s will be considered timely. the mailing date of this communication. D (35 U.S.C. § 133).		
Status					
1)⊠	Responsive to communication(s) filed on <u>07 O</u>	ctober 2004.	· ·		
2a) <u></u>		action is non-final.			
3)□	Since this application is in condition for allowar closed in accordance with the practice under E				
Disposit	ion of Claims				
4)⊠	Claim(s) 31 and 34 is/are pending in the applic	eation.			
	4a) Of the above claim(s) is/are withdraw	wn from consideration.			
5)	Claim(s) is/are allowed.				
6)⊠	Claim(s) 31 and 34 is/are rejected.				
7)	Claim(s) is/are objected to.				
8)[Claim(s) are subject to restriction and/or	r election requirement.	·		
Applicat	ion Papers				
9)	The specification is objected to by the Examine	r.			
·	10) The drawing(s) filed on is/are: a) accepted or b) objected to by the Examiner.				
	Applicant may not request that any objection to the	drawing(s) be held in abeyance. See	e 37 CFR 1.85(a).		
	Replacement drawing sheet(s) including the correct	ion is required if the drawing(s) is obj	jected to. See 37 CFR 1.121(d).		
11)	The oath or declaration is objected to by the Ex	aminer. Note the attached Office	Action or form PTO-152.		
Priority ı	under 35 U.S.C. § 119				
· <u>-</u>	•	priority under 25 H.S.C. \$ 440(a)	(d) or (5)		
	Acknowledgment is made of a claim for foreign ☐ All b)☐ Some * c)☐ None of:	priority under 35 U.S.C. 9 119(a)	i-(d) or (i).		
a):	1.☐ Certified copies of the priority documents	s have been received			
	Certified copies of the priority documents		on No		
	$3. \square$ Copies of the certified copies of the prior	rity documents have been receive			
* 5	application from the International Bureau See the attached detailed Office action for a list		.d		
	see the attached detailed Office action for a list	or the certified copies hot receive	:u.		
Attachmen	t(s)				
	e of References Cited (PTO-892)	4) Interview Summary			
	e of Draftsperson's Patent Drawing Review (PTO-948) mation Disclosure Statement(s) (PTO-1449 or PTO/SB/08)	Paper No(s)/Mail Da 5) Notice of Informat P	ate atent Application (PTO-152)		
	r No(s)/Mail Date	6) Other:			

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DETAILED ACTION

1. Applicant's response to the Office Actions mailed August 4, 2004 and October 7, 2004, on October 26, 2004, is acknowledged.

Claim Disposition

2. Claims 1-30, 32-33 and 35 have been canceled. Claims 31 and 34 are pending and under examination.

Claim Objection

3. Claim 31 is objected to because of the following informalities:

For clarity and precision of claim language it is suggested that the claim is amended to recite:

An assay method for identifying a compound able to modulate the interaction between (A) ATM (Ataxia-telangiectasia mutated) or ATR (ATM-Rad3-related) and (B) p53, the method including the steps of:

- (a) bringing into contact (1)ATM or ATR or a fragment of ATM or ATR which phosphorylates p53, (2) p53 or a fragment of p53 which includes a site which is phosphorylated by ATM or ATR and (3) a test compound; and
 - (b) determining phosphorylation at said site,

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wherein an increase or decrease in the phosphorylation at said site in the presence relative to the absence of the test compound being indicative that the compound is able to modulate the interaction between ATM or ATR and p53.

Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

- (a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.
- 4. This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. 103 (a), the examiner presumes that the subject matter of the various claims was commonly owned at the time any inventions covered therein were made absent any evidence to the contrary. Applicant is advised of the obligation under 37 CFR 1.56 to point out the inventor and invention dates of each claim that was not commonly owned at the time a later invention was made in order for the examiner to consider the applicability of 35 U.S.C. 103 (c) and potential 35 U.S.C. 102 (f) or (g) prior art under 35 U.S.C. 103 (a).

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5. Claims 31 and 34 are rejected under 35 U.S.C. 103 (a) as being unpatentable over Hoekstra et al. (WO 97/18323, May 22, 1997) in view of Meyn (Cancer Research, vol. 55, pages 5991-6001, 1995).

Hoekstra et al. teach an assay for identifying modulators of ATM (identified as the cell cycle checkpoint phosphatidylinositol kinase related protein) and MCSS1 gene, which is similar to p53 that involves contact with a test compound and a determination/quantification step (claim 31; see abstract, page 11 and claim 26 of the reference). Hoekstra et al. state that if a particular form of cancer results from a mutation in a gene such as p53, an agent which inhibits the transcription or the enzymatic activity may be used to render cancerous cells more sensitive to chemotherapy or radiation therapy (page 11). In-so-far-as Hoekstra et al. does not explicitly teach the modulation of the interaction between ATM and p53, Meyn teach that ATM physically interactions with p53 (page 5998) and suggests that ATM can phosphorylate p53 (claim 31). As it is well known in the art that ATM is part of a pathway that responds to DNA damage from ionizing radiation, thus ATM selectively regulates distinct p53-dependent cell cycle checkpoint and apoptotic pathways, a compound that modulates ATM will affect the interaction of ATM with p53 and modulate phosphorylation of p53 by ATM (claim 34).

Therefore, it would have been obvious to one of ordinary skill in the art at the time the invention was made to modify the teachings of Hoekstra et al. and include that modulation of ATM affects the interaction between ATM and p53 because Meyn state that ATM is upstream of p53 and that ATM and p53 physically interact (page 5998), thus inhibiting ATM would therefore affect the interaction between p53 and ATM. In addition, Meyn suggests that ATM, which acts similarly to DNA-PKcs, can phosphorylate p53 (page 5997). The skilled artisan

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would therefore, be motivated to add in the effect that modulating ATM affects the interaction of ATM with p53, because Hoekstra et al. disclose an assay method to identify compounds that modulate ATM and agents that would modulate genes such as p53, stating that the therapeutic value in such an agent lies in the fact that current radiation therapy or chemotherapy does nothing to overcome the ability of the p53 mutant cancerous cell to sense and correct the DNA damage imposed as a result of the treatment. Furthermore, it is known in the art that p53 induction is significantly reduced following exposure to ionizing radiation in AT cell lines, which demonstrates the need for said compound.

Thus, the claimed invention was obvious to make and use at the time it was made and was *prima facie* obvious.

6. Claims 31 and 34 are rejected under 35 U.S.C. 103 (a) as being unpatentable over Hoekstra et al. (WO 97/18323, May 22, 1997) in view of Baskaran et al. (Letters to Nature, vol. 387, pages 516-519, 1997).

Hoekstra et al. teach an assay for identifying modulators of ATM (identified as the cell cycle checkpoint phosphatidylinositol kinase related protein) and MCSS1 gene, which is similar to p53 that involves contact with a test compound and a determination/quantification step (claim 31; see abstract, page 11 and claim 26 of the reference). Hoekstra et al. state that if a particular form of cancer results from a mutation in a gene such as p53, an agent which inhibits the transcription or the enzymatic activity may be used to render cancerous cells more sensitive to chemotherapy or radiation therapy (page 11). In-so-far-as Hoekstra et al. does not explicitly teach the modulation of the interaction between ATM and p53, Baskaran et al. teach that ATM

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phosphorylates c-Abl and other proteins and that c-Abl and p53 are downstream targets of ATM, thus suggests that ATM can phosphorylate p53 (claim 31, see page 517 of the reference). As it is well known in the art that ATM is part of a pathway that responds to DNA damage from ionizing radiation, thus ATM selectively regulates distinct p53-dependent cell cycle checkpoint and apoptotic pathways, a compound that modulates ATM will affect the interaction of ATM with p53 and modulate phosphorylation of p53 by ATM (claim 34).

Therefore, it would have been obvious to one of ordinary skill in the art at the time the invention was made to modify the teachings of Hoekstra et al. and include that modulation of ATM affects the interaction between ATM and p53 because Baskaran et al. state that ATM is upstream of p53 and that ATM and p53 interact (page 517), thus inhibiting ATM would therefore affect the interaction between p53 and ATM. In addition, Baskaran et al. suggests that ATM phosphorylates p53 as it phosphorylates c-Abl and other proteins. The skilled artisan would therefore, be motivated to add in the effect that modulating ATM affects the interaction of ATM with p53, because Hoekstra et al. disclose an assay method to identify compounds that modulate ATM and agents that would modulate genes such as p53, stating that the therapeutic value in such an agent lies in the fact that current radiation therapy or chemotherapy does nothing to overcome the ability of the p53 mutant cancerous cell to sense and correct the DNA damage imposed as a result of the treatment. Furthermore, it is known in the art that p53 induction is significantly reduced following exposure to ionizing radiation in AT cell lines, which demonstrates the need for said compound.

Thus, the claimed invention was obvious to make and use at the time it was made and was *prima facie* obvious.

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7. The responses filed on August 4, 2004 and October 7, 2004 have been considered. The rejections of record have been withdrawn. The responses filed indicates that all claims are in condition for allowance, however, note that new grounds of rejections have been instituted as the prior art disclosed above reads on the present claims.

Conclusion

8. No claims are presently allowable.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Hope A. Robinson whose telephone number is 571-272-0957. The examiner can normally be reached on Monday-Friday.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Jon P. Weber, can be reached at (571) 272-0925. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

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Hope A. Robinson, MS-

Patent Examiner

JON WEBER

SUPERVISORY PATENT EXAMINER